69. A method according to claim 68, wherein detecting the label provides an indication of where the prostate cells are localized within the body of the human subject.

70. A method according to claim 69, wherein the label is detected using an imaging device.

71. A method according to claim 68, wherein the administering is carried out parenterally.

72. A method according to claim 71, wherein the administering is carried out intravenously.

73. A method according to claim 68, wherein the administering is carried out by intracavitary instillation.

74. A method according to claim 68, wherein the administering is carried out rectally.

75. A method according to claim 68, wherein the label is detected using a transrectal probe.

76. A method according to claim 68, wherein the antibody or antigen binding portion thereof is administered following a prostatectomy.

77. A method according to claim 68, wherein the antibody or antigen binding portion thereof is in a composition further comprising a pharmaceutically acceptable carrier, excipient, or stabilizer.

78. A method according to claim 68, wherein the antibody or antigen binding portion binds live cells and/or wherein the antibody is an IgG.

79. A method according to claim 68, wherein the antibody is selected from the group consisting of a monoclonal antibody and a polyclonal antibody.

80. A method according to claim 79, wherein the antibody is selected from the

group consisting of an E99, a J415, a J533, and a J591 monoclonal antibody.

81. A method according to claim 79, wherein the antibody is a monoclonal antibody produced by a hybridoma having an ATCC Accession Number selected from the group consisting of HB-12101, HB-12109, HB-12127, and HB-12126.

82. A method according to claim 68, wherein the antibody or antigen binding portion thereof binds to an epitope of prostate specific membrane antigen which is also recognized by a monoclonal antibody selected from the group consisting of an E99, a J415, a J533, and a J591 monoclonal antibody.

83. A method according to claim 82, wherein the antibody or antigen binding portion thereof binds to an epitope of prostate specific membrane antigen which is also recognized by monoclonal antibody J591.

84. A method according to claim 68, wherein the antibody or antigen binding portion thereof comprises an antigen binding portion of an amino acid sequence selected from the group consisting of SEQ ID NO:8 (variable heavy chain), SEQ ID NO:19 (variable light chain), an amino acid sequence of the variable heavy chain produced by the hybridoma having ATCC deposit no. HB-12126, and an amino acid sequence of the variable light chain produced by the hybridoma having ATCC deposit no. HB-12126.

85. A method according to claim 84, wherein the antibody or antigen binding portion thereof comprises an antigen binding portion of an amino acid sequence of SEQ ID NO:8 (variable heavy chain) or an amino acid sequence of the variable heavy chain produced by the hybridoma having ATCC deposit no. HB-12126 and an antigen binding portion of an amino acid sequence of SEQ ID NO:19 (variable light chain) or an amino acid sequence of the variable light chain produced by the hybridoma having ATCC deposit no. HB-12126.

86. A method according to claim 84, wherein the antibody or antigen binding portion thereof comprises an antigen binding portion of an amino acid sequence selected from the group consisting of SEQ ID NO:8 (variable heavy chain) and SEQ ID NO:19 (variable light chain).

87. A method according to claim 84, wherein the antibody or antigen binding portion thereof comprises an antigen binding portion of an amino acid sequence from SEQ ID NO:8 (variable heavy chain) and an antigen binding portion of an amino acid sequence from SEQ ID NO:19 (variable light chain).

88. A method according to claim 84, wherein the antibody or antigen binding portion thereof comprises an antigen binding portion of an amino acid sequence selected from the group consisting of an amino acid sequence of the variable heavy chain produced by the hybridoma having ATCC deposit no. HB-12126, and an amino acid sequence of the variable light chain produced by the hybridoma having ATCC deposit no. HB-12126.

89. A method according to claim 84, wherein the antibody or antigen binding portion thereof comprises an antigen binding portion of an amino acid sequence of the variable heavy chain produced by the hybridoma having ATCC deposit no. HB-12126 and an antigen binding portion of an amino acid sequence of the variable heavy chain produced by the hybridoma having ATCC deposit no. HB-12126.

90. A method according to claim 68, wherein the antibody or antigen binding portion thereof comprises an antigen binding portion of an amino acid sequence encoded by a nucleic acid sequence selected from the group consisting of SEQ ID NO:6 (variable heavy chain), SEQ ID NO:17 (variable light chain), a nucleic acid sequence which encodes the variable heavy chain produced by the hybridoma having ATCC deposit no. HB-12126, and a nucleic acid sequence which encodes the variable light chain produced by the hybridoma having ATCC deposit no. HB-12126.

91. A method according to claim 90, wherein the antibody or antigen binding portion thereof comprises an antigen binding portion encoded by a nucleic acid sequence of SEQ ID NO:6 (variable heavy chain) or a nucleic acid sequence which encodes the variable heavy chain of the hybridoma having ATCC deposit no. HB-12126 and an antigen binding portion encoded by a nucleic acid sequence of SEQ ID NO:17 (variable light chain) or a nucleic acid sequence which encodes the variable light chain produced by the hybridoma having ATCC deposit no. HB-12126.

92. A method according to claim 241, wherein the antibody or antigen binding portion thereof which comprises an antigen binding portion of an amino acid sequence encoded by a nucleic acid sequence selected from the group consisting of SEQ ID NO:6 (variable heavy chain) and SEQ ID NO:17 (variable light chain).

93. A method according to claim 90, wherein the antibody or antigen binding portion thereof which comprises an antigen binding portion of an amino acid sequence encoded by a nucleic acid sequence from SEQ ID NO:6 (variable heavy chain) and an antigen binding portion of an amino acid sequence encoded by a nucleic acid sequence from SEQ ID NO:17 (variable light chain).

94. A method according to claim 90, wherein the antibody or antigen binding portion thereof which comprises an antigen binding portion of an amino acid sequence encoded by a nucleic acid sequence selected from the group consisting of a nucleic acid sequence which encodes the variable heavy chain produced by the hybridoma having ATCC deposit no. HB-12126, and a nucleic acid sequence which encodes the variable light chain produced by the hybridoma having ATCC deposit no. HB-12126.

95. A method according to claim 90, wherein the antibody or antigen binding portion thereof which comprises an antigen binding portion of an amino acid sequence encoded by a nucleic acid which encodes the variable heavy chain produced by the

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hybridoma having ATCC deposit no. HB-12126 and an antigen binding portion of an amino acid sequence encoded by a nucleic acid which encodes the variable heavy chain produced by the hybridoma having ATCC deposit no. HB-12126.

96. A method according to claim 68, 78, 82, 84, or 90, wherein the antibody is a monoclonal antibody.

97. A method according to claim 68, 78, 82, 84, or 90, wherein the antibody or antigen binding portion thereof is internalized with the prostate specific membrane antigen.

98. A method according to claim 68, 78, 82, 84, or 90, wherein the antibody or antigen binding portion thereof is selected from the group consisting of a Fab fragment, a $F(ab')_2$ fragment, and a Fv fragment.

99. A method according to claim 68, 78, 82, 84, or 90, wherein the label is selected from the group consisting of a fluorescent label, a biologically-active enzyme label, a radiolabel, a nuclear magnetic resonance active label, a luminescent label, and a chromophore label.

100. A method according to claim 99, wherein the label is a radiolabel.

101. A method according to claim 100, wherein the radiolabel is a short-range radiation emitter.

102. A method according to claim 101, wherein the radiolabel is selected from the group consisting of ^{212}Bi, ^{213}Bi, and ^{211}At.

103. A method according to claim 100, wherein the radiolabel is selected from the group consisting of ^{32}P, ^{125}I, ^{3}H, ^{14}C, and ^{188}Rh.

104. A method according to claim 100, wherein the radiolabel is ^{131}I.

105. A method according to claim 100, wherein the radiolabel is ^{99}mTc.

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106. A method according to claim 100, wherein the radiolabel is ^{111}In.

107. A method according to claim 68, wherein the prostate cells are prostate epithelial cells. --

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